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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/572,316	03/17/2006	Robert Jongejan	27233U	8025
	7590 12/24/200 OCIATES PLLC	EXAMINER		
112 South West Street			BLIZZARD, CHRISTOPHER JAMES	
Alexandria, VA 22314			ART UNIT	PAPER NUMBER
			4185	
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Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)			
	10/572,316	JONGEJAN ET AL.			
Office Action Summary	Examiner	Art Unit			
	CHRISTOPHER BLIZZARD	4185			
The MAILING DATE of this communication ap Period for Reply	ppears on the cover sheet with the c	correspondence address			
A SHORTENED STATUTORY PERIOD FOR REPL WHICHEVER IS LONGER, FROM THE MAILING ID. - Extensions of time may be available under the provisions of 37 CFR 1 after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period. - Failure to reply within the set or extended period for reply will, by statu Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	DATE OF THIS COMMUNICATION .136(a). In no event, however, may a reply be tind d will apply and will expire SIX (6) MONTHS from te, cause the application to become ABANDONE	N. nely filed the mailing date of this communication. D (35 U.S.C. § 133).			
Status					
Responsive to communication(s) filed on 3/12 2a) This action is FINAL . 2b) This action is FINAL . 3) Since this application is in condition for allowed closed in accordance with the practice under	is action is non-final. ance except for formal matters, pro				
Disposition of Claims					
4) Claim(s) 1-25 is/are pending in the application 4a) Of the above claim(s) is/are withdra 5) Claim(s) is/are allowed. 6) Claim(s) 1-25 is/are rejected. 7) Claim(s) is/are objected to. 8) Claim(s) are subject to restriction and/ Application Papers 9) The specification is objected to by the Examin 10) The drawing(s) filed on 17 March 2006 is/are: Applicant may not request that any objection to the Replacement drawing sheet(s) including the correction.	awn from consideration. for election requirement. her. a) ☑ accepted or b) ☐ objected to the drawing(s) be held in abeyance. Section is required if the drawing(s) is objected to the	e 37 CFR 1.85(a). jected to. See 37 CFR 1.121(d).			
11) The oath or declaration is objected to by the E	xaminer. Note the attached Office	Action or form PTO-152.			
Priority under 35 U.S.C. § 119 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of: 1. Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received.					
Attachment(s) 1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date 6/16/2006.	4) Interview Summary Paper No(s)/Mail Da 5) Notice of Informal F 6) Other:	ate			

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DETAILED ACTION

Claim Rejections - 35 USC § 102

1. The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless -

- (b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.
- 2. Claims 1-5, 7, 8, 10, 13-22 and 25 are rejected under 35 U.S.C. 102(b) as being anticipated by Wolf (US 5,809,997).

Regarding claim 1, Wolf discloses a compliance monitor for a drug delivery device for administering a drug, comprising; a switch, in the form of a strain gauge dynamic sensing arm (1555), actuatable by a user on delivering a dose from the device (Abstract); a sensor (1560) for detecting whether the device is properly positioned in contact with or relative to the user's body for delivery of the dose (column 19, lines 21-41); and a processor (1540) coupled to the switch (1555) and the sensor (1560) (fig. 16) for recording whether or not the device was properly positioned when the switch was actuated.

Regarding claim 2, Wolf discloses a compliance monitor which does not affect the normal operation of the drug delivery device (column 3, lines 24-27).

Regarding claim 3, Wolf discloses a compliance monitor (1200)) which is removably attachable to the drug delivery device(1210) (column 17, lines 7-16).

Regarding claim 4, Wolf discloses a compliance monitor comprising a clock coupled to a processor (605) and in which the time of actuation of the switch is recorded (column 6, lines 25-30).

Regarding claim 5, Wolf discloses a compliance monitor wherein proper positioning of the drug delivery device is positioning in contact with or relative to the user's mouth, nose or skin (column 15, lines 58-60).

Regarding claim 7, Wolf discloses a compliance monitor wherein the drug delivery device is for oral administration of the drug (column 15, lines 58-60).

Regarding claim 8, Wolf discloses a compliance monitor wherein the drug delivery device is an inhaler operated by the user depressing a pressurized canister (1590) containing the drug, and wherein the switch is a pressure-operated switch (1555) actuatable as the user depresses the canister (figs. 17b and 17c).

Regarding claim 10, Wolf discloses a compliance monitor in which the sensor (425) is a temperature sensor, in the form of a thermistor, for sensing body temperature (columns 15,16; lines 58-67, 1-5).

Regarding claim 13, Wolf discloses a compliance monitor according in which a change in an output of the sensor characteristic of correct use of the drug delivery device is used to determine whether the device was properly positioned when the dose was delivered (column 15, lines 55-65).

Regarding claim 14, Wolf discloses a compliance comprising an output (415) for downloading data to a docking station or a computer (fig. 10).

Regarding claim 15, Wolf discloses a compliance monitor in which the data comprises a compliance record of use of the drug delivery device, including a record of whether the sensor output indicates that the device was properly positioned on each occasion that a dose has been delivered (column 3, lines 58-69) (column 15, 58-60).

Regarding claim 16, Wolf discloses a docking station (2030) for use with a compliance monitor (fig. 20).

Regarding claim 17, Wolf disclose a compliance monitor with computer-readable medium carrying a computer program for programming a general purpose computer to receive and process data downloaded from a compliance monitor (column 6, lines41-50).

Regarding claim 18, Wolf discloses a compliance monitor with a drug delivery device (Abstract).

Regarding claim 19, Wolf discloses a method of using a compliance monitor to monitor use of a drug delivery device for administration of a drug, comprising the steps of: determining when a user operates the device to deliver a dose of the drug (column 1, lines 11-20); sensing whether the device is properly positioned in contact with or relative to the user's body when the dose is delivered (column 15, lines 55-60); and recording for each operation of the-device whether or not the device was properly positioned (column 16, lines 2-5).

Regarding claim 20, Wolf discloses a method comprising the step of determining and recording the time of each operation of the device (column 1, lines 16-17).

Regarding claim 21, Wolf discloses a method in which the drug delivery device

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is for oral administration of the drug and proper positioning of the device is proper positioning in the user's mouth (column 15, lines 58-60).

Regarding claim 22, Wolf discloses a method comprising a step of downloading

Regarding claim 22, Wolf discloses a method comprising a step of downloading recorded data from the compliance monitor to a docking station or a computer to allow a compliance record to be reviewed (fig. 20) (column 6, lines 41-45).

Regarding claim 25, Wolf discloses a compliance monitor wherein the drug delivery device is for oral administration by inhalation (column 15, 55-60).

Claim Rejections - 35 USC § 103

3. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

Claims 6 and 9 are rejected under 35 U.S.C. 103(a) as being unpatentable over Wolf (US 5,809,997) in view of Reinhold (US 7,073,499 B1).

Regarding claim 6, Wolf fails to disclose a compliance monitor wherein the drug delivery device is for topical administration of the drug.

However, Reinhold et al. discloses a drug delivery device, in the form of an inhaler, for topical administration of a drug (column 14, lines61-63).

4. It would have been obvious to one of ordinary skill in the art to modify the invention of Wolf to include a drug delivery device as taught by Reinhold since doing so

would allow the invention to be used by patients with a wider range of medication needs.

Regarding claim 9, Wolf fails to disclose a compliance monitor wherein the drug delivery device is selected from the group consisting of a dry powder inhaler, a pressurized metered dose inhaler and a nebuliser.

However, Reinhold teaches the similarities of drug delivery devices, including dry powder inhalers, pressurized metered dose inhalers and nebulisers (column 1, lines 25-28).

- 5. It would have been obvious to one of ordinary skill in the art to modify the invention of Wolf to use various drug delivery devices as taught by Reinhold since doing so would allow the invention to be used by patients with a wider range of medication needs.
- 6. Claims 11 and 12 are rejected under 35 U.S.C. 103(a) as being unpatentable over Wolf (US 5,809,997) in view of Trueba (US 6,684,880 B2).

Regarding claim 11, Wolf fails to disclose a compliance monitor in which a sensor is a light sensor for sensing when the sensor is covered.

However, Truedba discloses a compliance monitor with a light sensor for sensing when the sensor is covered (column 13, lines 28-31).

7. It would have been obvious to one of ordinary skill in the art to modify the invention of Wolf to include a light sensor as taught by Trueba since doing so would provide a simple way of determining if the invention was properly used column 13, lines 28-31).

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Regarding claim 12, Wolf fails to disclose a compliance monitor in which the sensor is a conductivity sensor for sensing body conductivity.

However, Reinhold discloses a compliance monitor with one or more sensors for determining proper use of device.

8. It would have been obvious to one of ordinary skill in the art to modify the invention of Wolf to include a sensor as taught by Reinhold, and for the sensor to be a conductivity sensor, since doing so would provide a simple way of determining if the invention was properly used.

Conclusion

The prior art made of record and not relied upon is considered pertinent to applicant's disclosure. The following references are cited for disclosing related limitations of the applicant's claimed and disclosed invention.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Christopher Blizzard whose telephone number is (571)-270-7138. The examiner can normally be reached on Monday-Thursday 7:30 AM - 6:00 PM EST.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Terrell McKinnon can be reached on (571) 272-4797.

12/18/2008

/C. B./

Examiner, Art Unit 4185

/Terrell L Mckinnon/

Supervisory Patent Examiner, Art Unit 4185